This Listing of Claims will replace all prior versions, and listings, of claims in the application:

LISTING OF CLAIMS

Claim 1 (canceled)

Claim 2 (currently amended): The method of Claim 1 A method for the prophylaxis or treatment of a hyperlipidemic condition or disorder in a subject, said method comprising administering a first amount of an apical sodium co-dependent bile acid transporter inhibitor and a second amount of an HMG Co-A reductase inhibitor, wherein the apical sodium co-dependent bile acid transporter inhibitor comprises

or a pharmaceutically acceptable salt, ester or prodrug thereof.

Claims 3-15 (canceled)

Claim 16 (currently amended): The method of Claim 1 2 wherein the HMG Co-A reductase inhibitor is selected from the group consisting of mevastatin, lovastatin, simvastatin, pravastatin, fluvastatin, cerivastatin, atorvastatin, ZD-4522 rosuvastatin, and the pharmaceutically acceptable salts, esters, conjugate acids, and prodrugs thereof.

Claim 17 (currently amended): The method of Claim 1 16 wherein the HMG Co-A reductase inhibitor is selected from the group consisting of atorvastatin, simvastatin, pravastatin, 2D-4522 rosuvastatin, and the pharmaceutically acceptable salts, esters, conjugate acids, and prodrugs thereof.

Claim 18 (currently amended): The method of Claim 1 16 wherein the HMG Co-A reductase inhibitor comprises mevastatin, or a pharmaceutically acceptable salt, ester or prodrug thereof.

Claim 19 (currently amended): The method of Claim 1 17 wherein the HMG Co-A reductase inhibitor comprises atorvastatin, or a pharmaceutically acceptable salt, ester or prodrug thereof.

Claim 20 (currently amended): The method of Claim 1 17 wherein the HMG Co-A reductase inhibitor comprises simvastatin, or a pharmaceutically acceptable salt, ester or prodrug thereof.

Claim 21 (currently amended): The method of Claim 1 17 wherein the HMG Co-A reductase inhibitor comprises pravastatin, or a pharmaceutically acceptable salt, ester or prodrug thereof.

Claim 22 (currently amended): The method of Claim 1 16 wherein the HMG Co-A reductase inhibitor comprises lovastatin, or a pharmaceutically acceptable salt, ester or prodrug thereof.

Claim 23 (currently amended): The method of Claim 4 16 wherein the HMG Co-A reductase inhibitor comprises cerivastatin, or a pharmaceutically acceptable salt, ester or prodrug thereof.

Claim 24 (currently amended): The method of Claim 1 16 wherein the HMG Co-A reductase inhibitor comprises fluvastatin, or a pharmaceutically acceptable salt, ester or prodrug thereof.

Claim 25 (currently amended): The method of Claim 4 17 wherein the HMG Co-A reductase inhibitor comprises ZD 4522 rosuvastatin, or a pharmaceutically acceptable salt, ester, conjugate acid, or prodrug thereof.

Claim 26 (currently amended): The method of Claim 1 2 wherein the HMG Co-A reductase inhibitor comprises NK-104 pitavastatin, or a pharmaceutically acceptable salt, ester, conjugate acid, or prodrug thereof.

Claims 27-28 (canceled)

Claim 29 (currently amended): The method of Claim 27 A method for the prophylaxis or treatment of a hyperlipidemic condition or disorder in a subject, said method comprising administering a first amount of an apical sodium co-dependent bile acid transporter inhibitor and a second amount of an HMG Co-A reductase inhibitor, wherein the apical sodium co-dependent bile acid transporter inhibitor comprises the racemate of

or a pharmaceutically acceptable salt, ester or prodrug thereof.

Claims 30-39 (canceled)

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Claim 40 (currently amended): The method of claim 28 2 wherein the apical sodium co-

dependent bile acid transporter inhibitor and the HMG Co-A reductase inhibitor are administered

in a sequential manner.

Claim 41 (currently amended): The method of claim 28 2 wherein the apical sodium co-

dependent bile acid transporter inhibitor and the HMG Co-A reductase inhibitor are administered

in a substantially simultaneous manner.

Claim 42 (currently amended): The method of claim 28 2 wherein the weight ratio of apical

sodium co-dependent bile acid transporter inhibitor to HMG Co-A reductase inhibitor

administered is between about 1:50 to about 3:1.

Claim 43 (currently amended): The method of claim 28 2 wherein said apical sodium co-

dependent bile acid transporter inhibitor is administered in a daily dose ranging from about 0.008

mg to about 100 mg, and said HMG Co-A reductase inhibitor is administered in a daily dose

ranging from about 0.05 mg to about 100 mg.

Claim 44 (currently amended): The method of claim 28 2 wherein said apical sodium co-

dependent bile acid transporter inhibitor is administered in a daily dose range from about 0.08

<u>0.008</u> mg to about 100 mg.

Claim 45 (currently amended): The method of claim 28 2 wherein the HMG Co-A reductase

inhibitor is administered in a daily dose range from about 0.05 mg to about 100 mg.

Claim 46 (cancelled)

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Claim 47 (currently amended): The composition of Claim 46 wherein the A composition comprising a first amount of an apical sodium co-dependent bile acid transporter inhibitor comprises comprising

or a pharmaceutically acceptable salt, ester or prodrug thereof;

a second amount of an HMG Co-A reductase inhibitor, or a pharmaceutically acceptable salt, ester, conjugate acid, or prodrug thereof; and

a pharmaceutically acceptable carrier;

wherein the first and second amounts of said inhibitors together comprise a therapeutically effective amount of said inhibitors.

Claims 48-60 (cancelled)

Claim 61 (currently amended): The composition of Claim 46 47 wherein the HMG Co-A reductase inhibitor is selected from the group consisting of mevastatin, lovastatin, simvastatin, pravastatin, fluvastatin, cerivastatin, atorvastatin, <u>ZD-4522 rosuvastatin</u>, <u>NK-104 pitavastatin</u>, and the pharmaceutically acceptable salts, esters, conjugate acids, and prodrugs thereof.

Claim 62 (currently amended): The composition of Claim 46 61 wherein the HMG Co-A reductase inhibitor is selected from the group consisting of atorvastatin, simvastatin, pravastatin,

ZD-4522 rosuvastatin, and the pharmaceutically acceptable salts, esters, conjugate acids, and prodrugs thereof.

Claim 63 (currently amended): The composition of Claim 46 61 wherein the HMG Co-A reductase inhibitor comprises mevastatin, or a pharmaceutically acceptable salt, ester or prodrug thereof.

Claim 64 (currently amended): The composition of Claim 46 62 wherein the HMG Co-A reductase inhibitor comprises atorvastatin, or a pharmaceutically acceptable salt, ester or prodrug thereof.

Claim 65 (currently amended): The composition of Claim 46 62 wherein the HMG Co-A reductase inhibitor comprises simvastatin, or a pharmaceutically acceptable salt, ester or prodrug thereof.

Claim 66 (currently amended): The composition of Claim 46 62 wherein the HMG Co-A reductase inhibitor comprises pravastatin, or a pharmaceutically acceptable salt, ester or prodrug thereof.

Claim 67 (currently amended): The composition of Claim 46 61 wherein the HMG Co-A reductase inhibitor comprises lovastatin, or a pharmaceutically acceptable salt, ester or prodrug thereof.

Claim 68 (currently amended): The composition of Claim 46 61 wherein the HMG Co-A reductase inhibitor comprises cerivastatin, or a pharmaceutically acceptable salt, ester or prodrug thereof.

Claim 69 (currently amended): The composition of Claim 46 61 wherein the HMG Co-A reductase inhibitor comprises fluvastatin, or a pharmaceutically acceptable salt, ester or prodrug thereof.

Claim 70 (currently amended): The composition of Claim 46 62 wherein the HMG Co-A reductase inhibitor comprises ZD-4522 rosuvastatin, or a pharmaceutically acceptable salt, ester, conjugate acid, or prodrug thereof.

Claim 71 (currently amended): The composition of Claim 46 61 wherein the HMG Co-A reductase inhibitor comprises NK-104 pitavastatin, or a pharmaceutically acceptable salt, ester, conjugate acid, or prodrug thereof.

Claim 72 (currently amended): The composition of Claim 46 wherein the A composition comprising a first amount of an apical sodium co-dependent bile acid transporter inhibitor comprises comprising the racemate of

or a pharmaceutically acceptable salt, ester or prodrug thereof; and

the <u>an</u> HMG Co-A reductase inhibitor is selected from the group consisting of mevastatin, lovastatin, simvastatin, pravastatin, fluvastatin, cerivastatin, atorvastatin, <u>ZD-4522</u> rosuvastatin, <u>NK-104</u> pitavastatin, and the pharmaceutically acceptable salts, esters, conjugate acids, and prodrugs thereof.

Claims 73-83 (canceled)

Claim 84 (currently amended): The composition of claim 73 47 wherein the weight ratio of apical sodium co-dependent bile acid transporter inhibitor to HMG Co-A reductase inhibitor is between about 1:50 to about 3:1.

Claim 85 (canceled)

Claim 86 (currently amended): A kit of Claim 85 A kit containing a first dosage form comprising an apical sodium co-dependent bile acid transporter inhibitor and a second dosage form comprising an HMG Co-A reductase inhibitor wherein the apical sodium co-dependent bile acid transporter inhibitor comprises the 4R,5R enantiomer of

or a pharmaceutically acceptable salt, ester or prodrug thereof.

Claim 87 (currently amended): A <u>The</u> kit of Claim 86 wherein the HMG Co-A reductase inhibitor is selected from the group consisting of mevastatin, lovastatin, simvastatin, pravastatin, fluvastatin, cerivastatin, atorvastatin, <u>ZD-4522</u> rosuvastatin, <u>NK-104</u> pitavastatin, and the pharmaceutically acceptable salts, esters, conjugate acids, and prodrugs thereof.

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Claim 88 (currently amended): A <u>The</u> kit of Claim 86 87 wherein the HMG Co-A reductase inhibitor is selected from the group consisting of atorvastatin, simvastatin, pravastatin, <u>ZD-4522 rosuvastatin</u>, and the pharmaceutically acceptable salts, esters, conjugate acids, and prodrugs thereof.

Claim 89 (canceled)